



	Dr. Vinay Chopra MD (Pathology & Micro Chairman & Consultant		icrobiology)	st l	Dr. Yugam MD CEO & Consultant	(Pathology	y)
NAME	: Mr. HARB	ANS SINGH					
AGE/ GENDER	: 85 YRS/MALE		PATIENT ID		: 1685131		
COLLECTED BY	:			REG. NO./LAB NO.		: 012411280060	
REFERRED BY	Y :			REGISTRATION DATE		: 28/Nov/2024 04:55 PM	
BARCODE NO.	:01521633			COLLECTION DATE		: 28/Nov/2024 05:29PM	
CLIENT CODE.	: KOS DIAGN			REPORTING DATE		: 28/Nov/2024 06:06PM	
CLIENT CODE. CLIENT ADDRESS						5V/2024 00.00FM	
CLIENI ADDRESS	: 0349/1, M	CHOLSON ROAD, AM	IDALA CANT I				
Test Name			Value		Unit		Biological Reference interval
HAEMATOLOGY							
GLYCOSYLATED HAEMOGLOBIN (HBA1C)							
GLYCOSYLATED HAEMOGLOBIN (HbA1c): WHOLE BLOOD			11.8 ^H %		4.0 - 6.4		
by HPLC (HIGH PERFORMANCE LIQUID CHR		· ·					
ESTIMATED AVERAGE PLASMA G			291.96 ^H		mg/dL		60.00 - 140.00
INTERPRETATION:							
AS PER AMERICAN DIABETES ASSOCIATION (ADA):							
REFERENCE GROUP GLYCOSYLATED HEMOGLOGIB (HBAIC) in %							%
Non diabetic Adults >= 18 years			<5.7				
At Risk (Prediabetes)			5.7 - 6.4				
Diagnosing Diabetes >= 6.5							
			Age > 19 Years Goals of Therapy:			< 7.0	
Therapeutic goals for glycemic control		emic control	Actions Suggested:			>8.0	
			Age < 19 Years				
			Goal of therapy:		<7.5		
2.Since Hb1c reflects lo concentration of HbAlc 3.Target goals of < 7.0 patients with significan appropiate.	ng term fluctua c. Converse is tr % may be bene t complications	tions in blood glucose ue for a diabetic previc ficial in patients with s of diabetes, limited lif	concentration, pusly under goo hort duration c e expectancy of	a diabetic od control of diabetes r extensive	patient who has rec but now poorly cont , long life expectanc e co-morbid conditio	ently unde rolled. y and no s ns, targett	regimen in diabetic patients. er good control may still have high ignificant cardiovascular disease. In ting a goal of < 7.0% may not be Ilar and nerve complications

KOS Diagnostic Lab

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5.Any condition that shorten RBC life span like acute blood loss, hemolytic anemia falsely lower HbA1c results.
6.HbA1c results from patients with HbSS, HbSC and HbD must be interpreted with caution, given the pathological processes including anemia, increased red cell turnover, and transfusion requirement that adversely impact HbA1c as a marker of long-term gycemic control.

7.Specimens from patients with polycythemia or post-splenctomy may exhibit increse in HbA1c values due to a somewhat longer life span of the red cells.

*** End Of Report ***



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